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Functionally graded ferroelectric polyetherimide composites for high temperature sensing

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High temperature ferroelectrics for thermally stable devices that can detect pressure and temperature are of great industrial interest. Here we describe composites of lead titanate (PT) particle-polyetherimide (PEI) polymers with stable dielectric and piezoelectric properties over a broad range of temperature and frequency. The reported materials have a low dielectric loss ($\tan \delta \sim 0.001$ at 1 kHz) and a high piezoelectric voltage coefficient of 100 mV m N⁻¹ at record temperatures of 175 °C. We demonstrate that a small ceramic loading leads to a significant change in thermally stable piezoelectric behavior, while the processability as well as mechanical properties remain comparable to those of the neat polymer. Careful design of the microstructure is performed by dielectrophoretic assembly of ferroelectric PT micro-particles to induce micro-wire configurations, which is shown to be a key element in attaining high functionality at low ceramic loading. Thermal imidization of the composites is performed in two steps, first partial imidization at 60 °C to form free standing films containing polyamic acid, followed by full imidization at 200 °C and 300 °C. The presence of highly polar polyamic acid results in higher dielectric permittivity and electrical conductivity that facilitate efficient poling. Upon complete imidization of the films at 300 °C the dielectric and piezoelectric properties are tested at elevated temperatures. A fully imidized composite contains completely closed imide groups, resulting in a thermally stable material with a very low dielectric loss that maintains more than 85% of its room temperature piezoelectric sensitivity up to 175 °C. The room temperature piezoelectric voltage coefficient shows more than 400% improvement over that of PT ceramics.

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1 Introduction

High temperature piezoelectric materials for application in automotive, aerospace and energy production have been a growing area of research.^{1–3} In automotive electronics, repeated thermal cycles up to 150 °C and temperatures as high as 500 °C are reported.⁴ In the aerospace industry, structural health monitoring of the engines requires sensors to be as close as possible to the engine, leading to exposure to high temperatures.^{5,6} Moreover, exploration of new sources for clean energy, such as nuclear and wind power, as well as geothermal and electrical energy, has significantly motivated the development of high temperature electronic materials.⁷ High temperature sensors for structure health monitoring can ensure enhanced lifetime and reliability,

less maintenance and downtime, and can thus reduce the overall cost of renewable energy sources.^{8,9} Piezoelectric materials can also be used to harvest mechanical energy from mechanical strain or vibrations present in the environment. An example of energy harvesting under extreme conditions is in oil producing wells, where the ambient pressures may reach 200 MPa, and the temperature reaches 200 °C. In this case local power production is crucial, since transmitting power from the surface is complicated, due to the difficulty in making reliable electrical connections in harsh environments.¹⁰

The use of piezopolymers in those applications is limited by their low operating temperatures. Polymer composites, however, have broader potential applications due to their excellent property tunability. Piezocomposites have been reported to be excellent sensing devices at room temperature offering a wide range of properties. However their usage in high temperature applications has been limited, due to the reduced sensing capability of piezo-ceramics at elevated temperatures, increased conductivity and mechanical attenuation, as well as variation of the piezoelectric properties of the ceramic phase with temperature.^{11,12} Ferroelectric materials get depolarized and lose piezoelectricity above their respective Curie temperatures (T_c). Moreover the low glass transition temperature and high thermal expansion of the

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polymer matrix limit the operational temperature range of piezocomposites.^{13,14} To overcome this problem piezoceramics with a high Curie temperature can be embedded in high temperature polymers. In practice, the operating temperature must be substantially below the Curie temperature of the piezoceramic phase, in order to minimize thermal aging and property degradation.

Amorphous aromatic polyimides, as a class of high-performance polymers, are interesting candidates for the applications mentioned above.^{15–19} Growing interest in viable, multifunctional and lightweight materials for demanding space applications has led to the development of a new series of such polymers^{15,17,20} with superior mechanical, chemical and electrical properties. Some of the polymer systems in this family have been reported to show insignificant piezoelectric properties.^{15,16} One of the most widely reported piezo polyimides is a polymer containing a single nitrile group, (β -CN)APB/ODPA. As opposed to the well-known semicrystalline PVDF polymer family, the high temperature performance of this polyimide allows these films to retain 50% of the room temperature remnant polarization at temperatures up to 150 °C.¹⁶ However, the remnant polarization of this system is very low, 14 mC m⁻², when poled at 100 kV mm⁻¹, for 1 h above T_g . Therefore, the piezoelectric charge coefficient of this polyimide is much lower than that of PVDF.^{15,16} Limited chain mobility in the imidized film restricts the preferential alignment of the dipoles upon application of an electric field. To solve this problem, new polymers have been synthesized, with different types and concentrations of the polar groups.¹⁸ Despite the interesting research outcomes, the polyimide systems exhibit very low piezoelectric responses that are not high enough to attract commercial interest.^{15,17,18,21}

The effect of processing parameters such as curing and poling conditions has also been investigated. The partially cured, corona poled polymers exhibited improved remanent polarization.²¹ In a more recent work *in situ* poling and imidization of partially cured (β -CN)APB/ODPA were investigated to maximize the degree of dipolar orientation.¹⁵ It has been shown that polyamic acid exhibits higher net dipole moment than the fully imidized polymer.^{15,17,18}

To overcome these limitations, and to achieve adequate combinations of mechanical, thermal and functional properties ferroelectric ceramic particles are embedded within a polyimide polymer. Structuring the ceramic particles into a chain form configuration, by means of dielectrophoresis (DEP), has been previously reported to improve the dielectric and piezoelectric properties of piezocomposites in various polymer systems.^{13,22,23} In this work, processing of high temperature lead titanate–polyetherimide composites is presented. Among various kinds of commercially available ferroelectric ceramics for sensing, lead titanate (PT) is chosen based on the high Curie temperature of 490 °C, large spontaneous polarization and a small relative dielectric constant.^{2,24,25} Therefore, it serves as a viable candidate for integration within high temperature polyimide films. Amongst high performance thermoplastics, PEI polymers composed of all rigid aromatic and heterocyclic monomers show high decomposition temperatures in excess of 450 °C, high glass

transition temperatures of above 200 °C, and outstanding mechanical properties (E of 3–4 GPa).^{26,27} The flexible ether bridge in the ODPA molecule can favorably influence the flexibility of the final polymer film.²⁸ Here we present a detailed study of the manufacturing procedure, microstructural analysis and functional properties of high temperature piezocomposites with DEP assembled PT particles in a 4,4'-oxydiphthalic anhydride (ODPA)/3,4'-oxydianiline diamine (ODA) polymer matrix. By combining experiment and simulations an understanding of the macroscopic properties and tuning strategies is provided.

2 Experimental procedure

2.1 Composite manufacturing

Polyetherimide polymers (PEI) are conventionally synthesized in two steps including precursor (polyamic acid (PAA)) formation followed by thermal imidization. The first step of preparation of PAA comprises the reaction of an aromatic diamine with an aromatic dianhydride in aprotic polar solvents. The scheme is shown in Fig. 1a. The PAA precursor in the current work was prepared *via* a solution of 4,4'-oxydiphthalic anhydride (ODPA)/3,4'-oxydianiline diamine (ODA) in *N*-methyl-2-pyrrolidone (NMP). The thermal properties of the films were determined by Thermogravimetric Analysis (TGA), using a Perkin Elmer Pyris Diamond, as well as Differential Scanning Calorimetry (DSC), using a Perkin Elmer Sapphire DSC. Samples were heated at a rate of 20 °C min⁻¹ under a nitrogen atmosphere to 550 °C. Dynamic Mechanical Thermal Analysis (DMTA) was performed in the tensile mode, at a frequency of 1 Hz and a heating rate of 2.0 °C min⁻¹, using films of approximate dimensions of 20 × 3 × 0.005 mm.

Lead titanate (PT) ceramic powder, purchased from Sigma Aldrich, was calcined at 800 °C for 2 h to develop single phase tetragonal PbTiO₃ as shown in Fig. 1a. The agglomerated powder was then dry-milled, using 5 mm zirconium balls for 2 h in a single G90 jar mill. The particle size distribution of milled powder in an aqueous solution with 10% isopropyl alcohol, measured using a Beckman Coulter LS230 laser diffraction analyzer was found to be $d(50) = 3$ μ m. The powder was stored in a vacuum drying oven at 120 °C for 24 h prior to the experiment to avoid moisture absorption. The dried PT powder was predispersed in NMP (Fig. 1a), stirred for 1 h, and added to polyamic-acid. The mixture was cast on an aluminum foil using a doctor blade, with an initial film thickness of 600 nm (Fig. 1b). Structuring of the particles in the polyamic-acid polymer was realized by a dielectrophoresis process, in which an electric field of 2 kV mm⁻¹ and a frequency of 2 kHz was applied to the composite medium of particles dispersed in PAA, using a function generator (Agilent, 33210A) coupled to a high voltage amplifier (Radiant Technologies Inc., T6000HVA-2) at room temperature (RT) for 1 h in an Argon purged oven as shown in Fig. 1b. Once the structuring of the particles was realized, as shown in Fig. 1b, the film was dried at 60 °C for 1.5 h under the same atmosphere and electric field. The randomly dispersed samples were obtained by the process described above, only

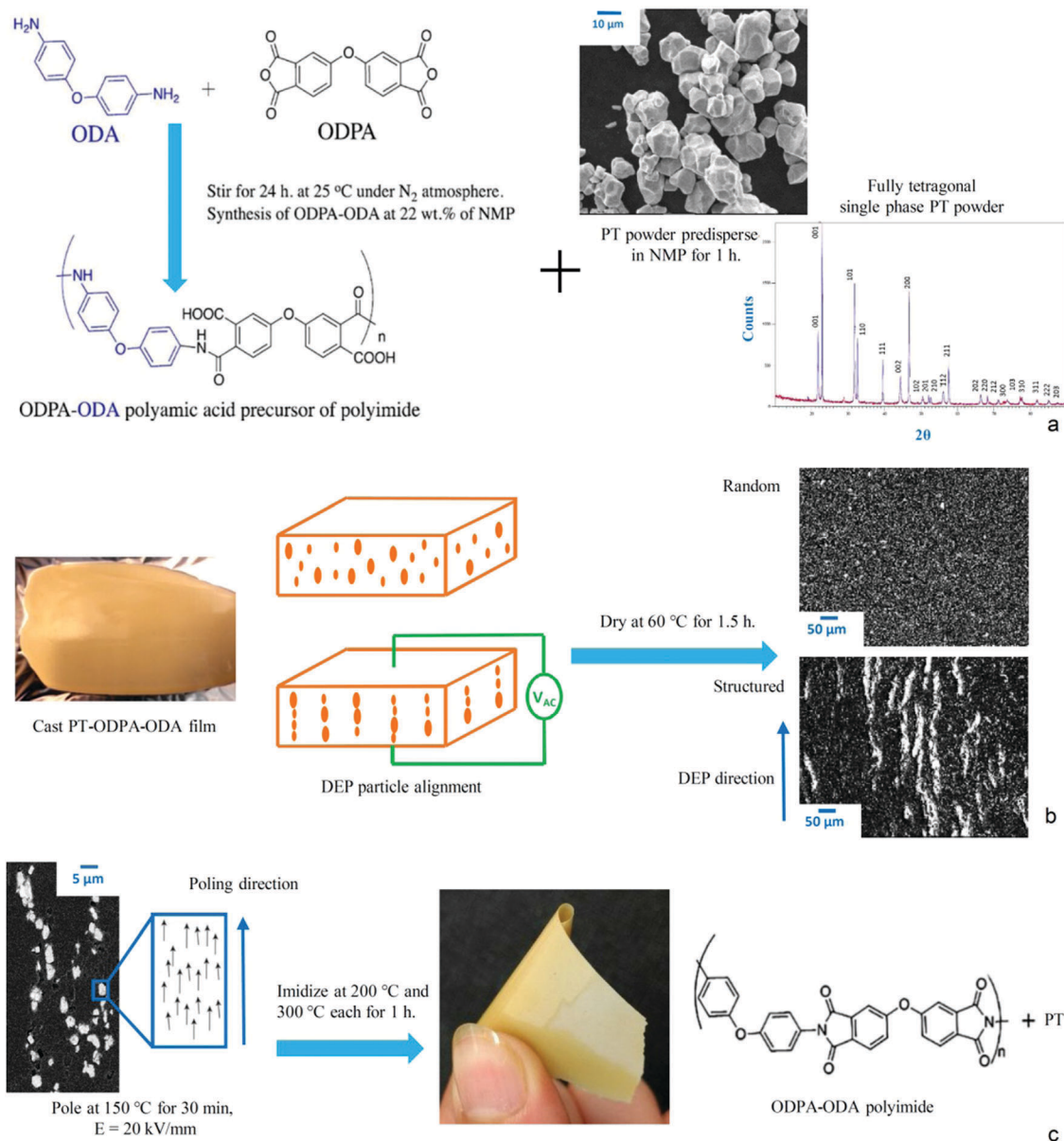


Fig. 1 Production route of PT-ODPA-ODA composites. (a) Scheme of ODPA-ODA polymerization. (b) Dielectrophoresis structuring of particles. (c) Poling and imidization.

without applying an electric field by oven curing at 60 °C for 1.5 h, using inert Argon gas as the surrounding medium. The free standing films were then removed and cut into discs of 7 mm diameter. Au electrodes were sputtered on the samples, and they were poled at 150 °C for 30 min with an electric field of 20 kV mm⁻¹. The poled films were then imidized in a vacuum oven at 200 °C and 300 °C, each for 1 h. The resulting flexible films have a fully imidized chemical structure as shown in Fig. 1c.

2.2 Measurement procedure

Novocontrol Broadband Dielectric Spectroscopy (BDS) is used in conjunction with a Cryostat high temperature sample cell and an Agilent E4991A impedance analyzer at 1 V and

1 kHz to measure the dielectric properties as well as electrical conductivity of the polymer films. The dielectric constant of the composites was determined by the parallel plate capacitor method using an Agilent 4263B LCR meter at 1 V and 1 kHz. The piezoelectric coefficient, d_{33} , measurements were done using a Berlincourt type M3001 d_{33} meter, KCF technologies (State College, PA) at 110 Hz. Before performing the d_{33} measurements, the samples were grounded with their electrodes shorted at 100 °C for 24 h to remove any remaining space charges. To test the d_{33} values of the composites at elevated temperatures, a modified sample holder was manufactured and connected to the CF technologies PM3001 d_{33} meter, as shown in Fig. 2. The sample was placed on a copper electrode and heated uniformly using two integrated programmable heaters.

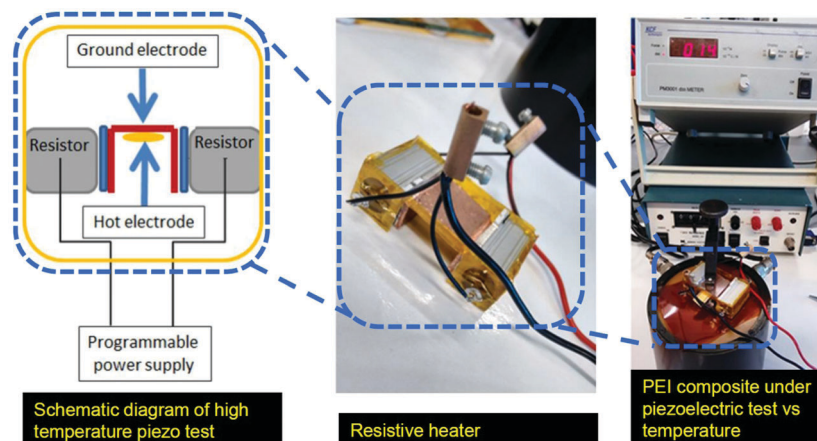


Fig. 2 The set-up platform for high temperature piezoelectric test showing the sample holder used in conjunction with Berlincourt M3001 for d_{33} measurements at elevated temperatures.

The fixture was thermally insulated using Kapton foils, and the set-up was calibrated using reference PZT ceramics with known properties. It has to be noted that the maximum test temperature is above the advised operating temperature of the set-up. Static uniaxial tensile tests were performed using an Instron 3369 tensile testing machine with a 50 N static load cell (62 291) with a speed of 0.1 mm min^{-1} at 20°C . Five samples having approximate dimensions of $40 \times 6 \times 0.045$ were tested and their average properties were calculated.

3 Results and discussion

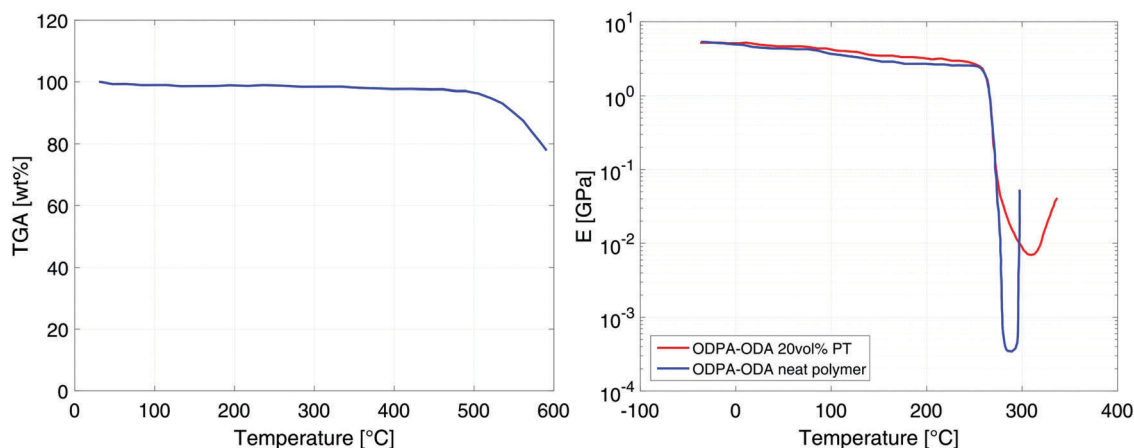
3.1 Microstructure analysis

The effect of DEP structuring on the SEM microstructures of the 20% PT-ODPA/ODA composite samples is shown in Fig. 1b and c. PT particles are homogeneously dispersed in the random composite, and no sign of sedimentation is observed. It can be seen that during dielectrophoresis, ceramic particles form

chain-like structures along the applied electric field direction. The chain-like particle assembly takes place as a result of attractive forces in the regions of the highest electric field gradient, as we have shown before computationally.²⁹ The degree of alignment which affects the final dielectric and electroactive properties of the composites depends on the force acting on the particles, as a consequence of the imposed electric field.

3.2 Thermo-mechanical analysis

Fig. 3a shows the thermogravimetric behavior of the obtained PEI material. An excellent thermal stability is observed and 2% weight loss is recorded at 520°C . The DMTA results, presented in Fig. 3b, show a storage modulus of 4.0 GPa and 5.0 GPa, at room temperature for neat ODPA/ODA and ODPA/ODA with 20 vol% PT, respectively. The T_g remains unchanged at 265°C which is in good agreement with the T_g of similarly reported compositions.^{30,31} The increase in high temperature storage modulus of the composite can be attributed to the enhancement of interaction between the polyetherimide matrix and the



(a) TGA spectrum of fully imidized ODPA/ODA.

(b) DMTA results of ODPA/ODA and 20% PT-ODPA/ODA.

Fig. 3 Thermo-mechanical properties of fully imidized PT-ODPA/ODA.

PT particles, which limits the segmental movement of the polymer.³²

3.3 Dielectric and piezoelectric analysis

The dependence of dielectric permittivity and electrical resistivity of fully imidized ODPA/ODA (PEI), as well as partially imidized ODPA/ODA (PAA) on frequency and temperatures is presented in Fig. 4. The permittivity of all the samples decreases with frequency (Fig. 4a and b), but the decreasing degree of the partially imidized ODPA/ODA is much higher than that of the fully imidized ODPA/ODA as shown by Ding *et al.*³³

For the fully imidized polymer at temperatures below T_g the polymer shows a stable dielectric behavior³⁴ with dielectric permittivity values ranging from 3.3 to 3.4 (Fig. 4a). Increasing the frequency from 1 MHz to 100 Hz results in a slight decrease in permittivity, which can be attributed to the dipolar dispersion. In other words dipoles gradually cannot follow the applied external field as the frequency increases.^{30,35,36} As the temperature approaches the glass transition temperature of fully imidized ODPA/ODA (260 °C) the dielectric permittivity increases due to the enhanced dipole motion.^{37,38} At low temperatures mainly electronic

and atomic polarizations contribute to the permittivity.³⁶ Similar behavior has been reported for (β -CN) APB/ODPA polyimides.²⁰

The partially imidized ODPA/ODA film shows a higher dielectric permittivity than the fully imidized ODPA/ODA film, as PAA possesses more polar groups ($-\text{COOH}$) than PEI, as shown in Fig. 1a. Furthermore a more significant frequency dependent permittivity behavior is observed for partially imidized ODPA/ODA (Fig. 4b). The increase of dielectric permittivity at low frequencies can be contributed to higher interface polarization.³⁹ The structures of dianhydrides have been shown to influence the dielectric properties, since ketone (benzophenone) groups are easily polarized in the electric field.⁴⁰

The resistivity plots (Fig. 4c and d) show that the electrical conductivity of partially imidized ODPA/ODA at 100 °C is more than two orders of magnitude higher than that of fully imidized ODPA/ODA at this temperature. This is in agreement with the results of Park *et al.*, who investigated the effect of processing parameters such as curing conditions on (β -CN) APB/ODPA systems.¹⁵ It has been shown that the partially cured (β -CN)APB/ODPA, in other words polyamic acid, exhibits a higher net dipole moment than the fully imidized polymer.^{15,17,18}

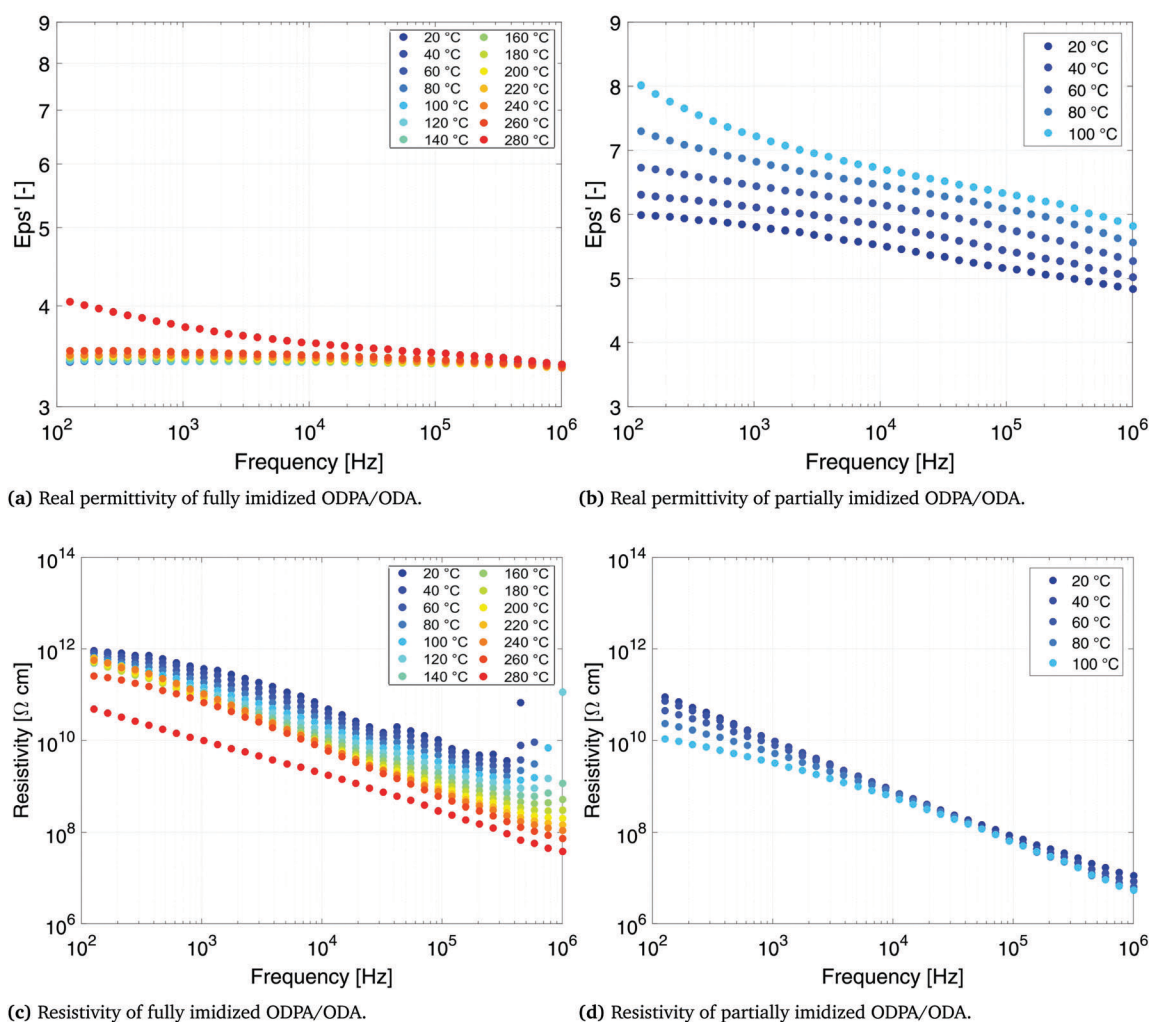


Fig. 4 Dielectric and electrical resistivity properties of fully and partially imidized ODPA/ODA.

The lower resistivity of the partially imidized ODPA/ODA (Fig. 4d) facilitates efficient poling of the embedded ferroelectric PT particles. During poling, the mobility of charges in the polymer matrix is a key parameter in defining the final poling efficiency. As the dielectric permittivity of PT is much higher than that of the polymer matrix⁴¹ the charge relaxation time is mainly governed by the electrical conductivity of the matrix. The higher conductivity of the partially imidized ODPA/ODA shortens the time required for the electric field acting on the ceramic inclusion to reach saturation, thereby making the poling of the ceramic phase more efficient.^{42,43} Moreover it eliminates the need to use conductive fillers such as single-wall carbon nanotubes (SWNTs) that have been reported to enhance the poling efficiency of (β -CN) APB/ODPA polyimides due to better dipolar alignment with the poling field.²⁰ $\tan \delta$, as an indication of dielectric loss angle, is 0.001 and 0.02 for fully imidized and partially imidized ODPA/ODA films at 20 °C and 1 kHz, respectively. Increasing either temperature or frequency would increase the dielectric loss, as expected and reported.³⁴ High electrical resistivity of fully imidized ODPA/ODA shows the good insulation performance of these materials.

The effect of DEP structuring on the dielectric constant of 10%, 20% and 30% PT-ODPA/ODA composites is shown in Fig. 5a. The experimental data for random composites are fitted to Yamada's model, using an n value of 5.⁴⁴ The composite is assumed to be a uniform distribution of ellipsoidal particles in an isotropic polymer matrix. The dielectric constant of the composite in the poling direction is given by

$$\varepsilon_{\text{random}} = \varepsilon_{\text{m}} \left(1 + \frac{n\phi(\varepsilon_{\text{c}} - \varepsilon_{\text{m}})}{n\varepsilon_{\text{m}} + (\varepsilon_{\text{c}} - \varepsilon_{\text{m}})(1 - \phi)} \right) \quad (1)$$

where ε is the dielectric constant, and ϕ is the volume fraction of the ceramic phase. The subscripts c, m, and random refer to the ceramic, polymer matrix and 0–3 composite properties, respectively, and n is the inverse of the depolarization factor for an ellipsoidal particle in the direction of the applied electric field.

Dielectric properties of the structured composites are fitted to Bowen's model²² as shown in Fig. 5a. According to this model, the composite is considered as a collection of particles aligned as chain-like structures along a specific direction separated by polymer gaps. The equation for the dielectric permittivity for such a composite is as follows:

$$\varepsilon_{\text{structured}} = \phi_{\text{c}} \left(\frac{R\varepsilon_{\text{c}}\varepsilon_{\text{m}}}{\varepsilon_{\text{c}} + R\varepsilon_{\text{m}}} \right) + (1 - \phi_{\text{c}})\varepsilon_{\text{m}} \quad (2)$$

where $\varepsilon_{\text{structured}}$ is the dielectric constant of dielectrophoretically structured 1–3 composites, and R is the ratio of average particle size to the effective interparticle distance, which influences the fraction of the applied electric field acting on the ceramic particles. The value of R obtained for the best fit of the experimental data to Bowen's model was 10.

The influence of DEP structuring on piezoelectric charge constant, d_{33} , of the composites is shown in Fig. 5b, where the experimental d_{33} values of random and structured 10%, 20% and 30% PT-ODPA/ODA composites are compared with the predictions of Yamada's and van den Ende's models.¹³ According to Yamada the piezoelectric charge constant, d_{33} , of the composite in the poling direction is given by

$$d_{33\text{random}} = \frac{\phi\alpha n\varepsilon_{\text{random}}d_{33\text{c}}}{n\varepsilon_{\text{random}} + \varepsilon_{\text{c}} - \varepsilon_{\text{random}}} \quad (3)$$

where α is the poling ratio of the ceramic particles. The other parameters were introduced earlier using eqn (1). The best fit of the experimental data to the model predictions was obtained for $n = 5$ and $\alpha = 1$, which is an indication of high poling efficiency attributed to the relatively low electrical resistivity of the partially imidized ODPA/ODA.

The model proposed by van den Ende *et al.*²³ is applied to the structured composites. This model is an extension of Bowen's model,²² in which the d_{33} values of composites are obtained by modeling the particle-matrix sequences in the chains as two capacitors in series in the electrical domain, and two springs in

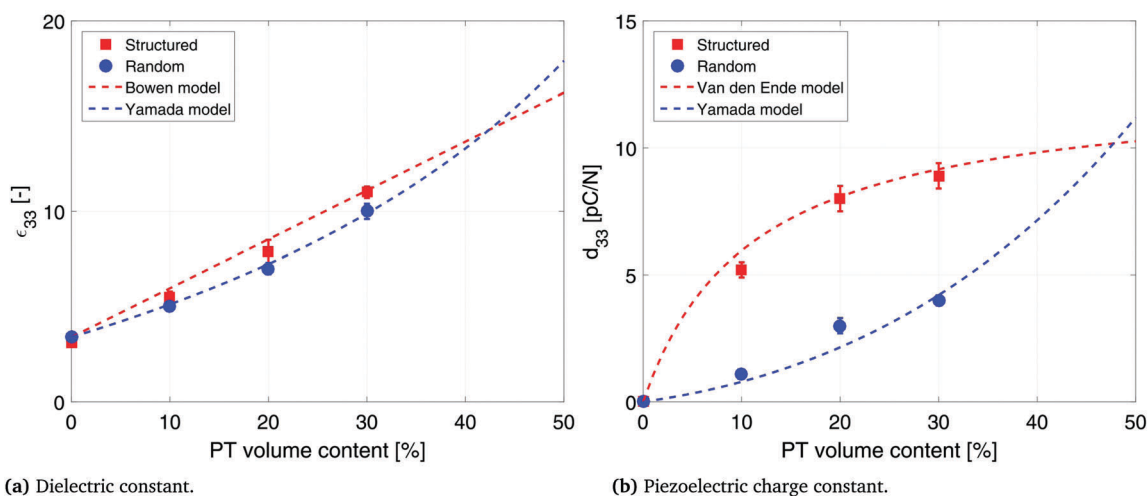


Fig. 5 Dielectric and piezoelectric properties of structured and random 10%, 20% and 30% PT-ODPA/ODA composites with their associated models.

series in the mechanical domain. The equation for the d_{33} values of the 1–3 composite is given by

$$d_{33\text{structured}} = \frac{(1 + R)^2 \epsilon_m \phi_c d_{33c} Y_c}{\epsilon_c + R \epsilon_m [(1 + R \phi_c) Y_c + (1 - \phi_c) R Y_m]} \quad (4)$$

where Y_m and Y_c are elastic moduli of the polymer matrix and that of the ceramic in the direction of chains, respectively, while R is the ratio of average particle size to the effective interparticle distance in the direction of the electric field. As shown in Fig. 5b, the values for both 0–3 and 1–3 composites match reasonably well to the models. The best fit of the model to the experimental data was obtained for an R of 10 which is in great agreement with the value of R obtained for the dielectric constant data.

The interfacial interaction between the ceramic particles and the polymer influences the final dielectric and piezoelectric properties of the composite materials.⁴⁵ The dielectric constant and piezoelectric charge constant of a composite are determined for understanding the nature of the interactions between the constituents of the composite. Lack of interfacial

adhesion between the ceramic particles and the polymeric matrix would result in a decrease in the dielectric constant and piezoelectric coefficients of the composite materials.^{46,47} Analytical models used in this work account for ceramic–polymer interaction as well as size and shape of the constituent particles. The good agreement between the experimental data and predictions of the model is an indication of good interfacial adhesion and pore free interfaces, which is confirmed by SEM microstructures (as shown in Fig. 1c). It is observed that PT particles are fully encapsulated by the polymer, while aligning in the direction of the applied electric field in structured composites. Thus interface porosity has not been observed in either of the random and structured composites.

The d_{33} values of 20% and 30% PT-ODPA/ODA composites are tested as a function of temperature, as shown in Fig. 6. A remarkably stable behavior is observed over the temperature range from room temperature to 175 °C. Furthermore, the room temperature d_{33} value can be retained upon cooling back. This stability is caused by the thermally stable dielectric and piezoelectric properties of the high T_c lead titanate,⁴⁸ as well as

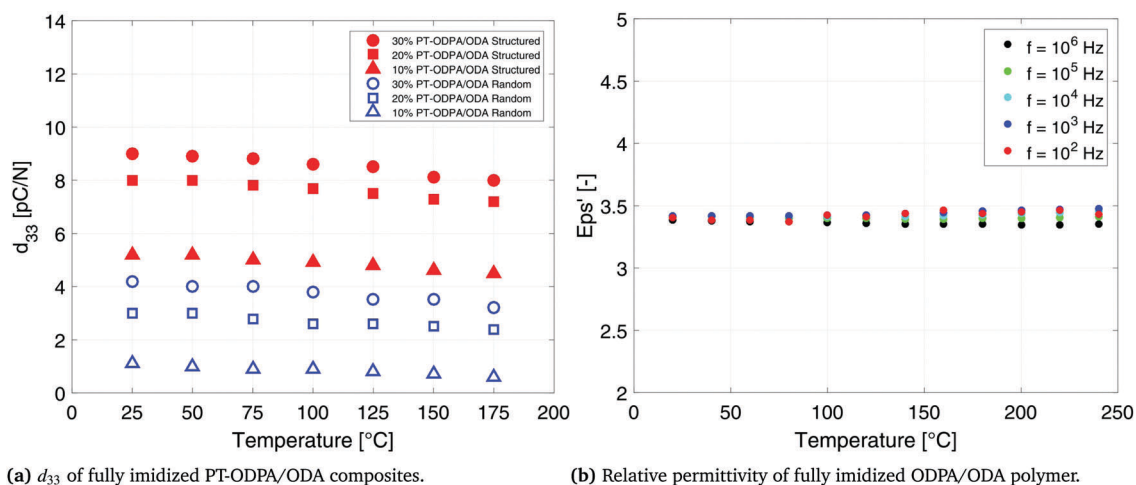


Fig. 6 Temperature dependent behavior of the ODPA/ODA polymer and 10%, 20% and 30% random and structured PT-ODPA/ODA composites.

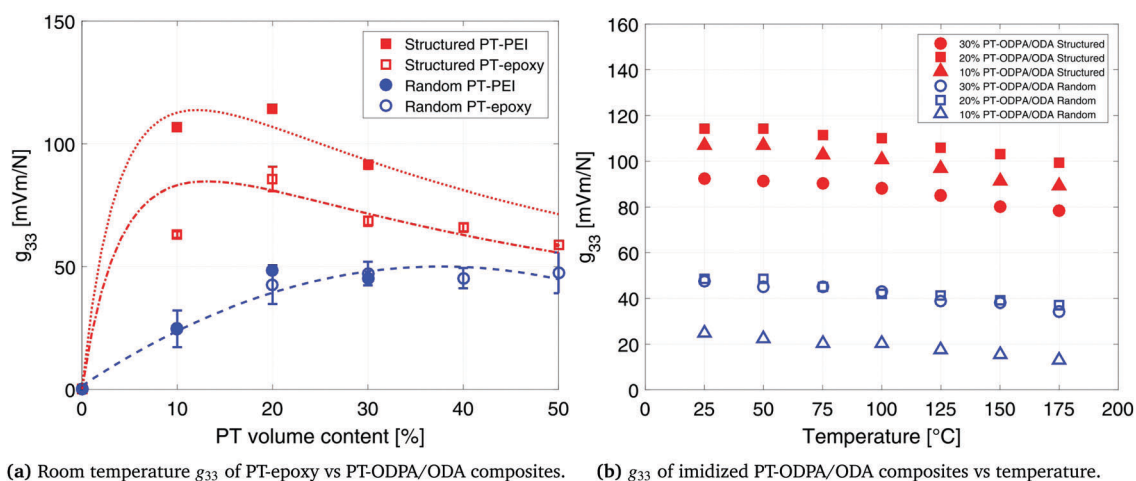


Fig. 7 g_{33} of 10%, 20% and 30% random and structured PT-ODPA/ODA composites.

stable dielectric properties of the ODPA/ODA composites in this temperature range, as shown in Fig. 6. Similar behavior has been reported for other PEI polymers.^{34,49}

The piezoelectric voltage coefficient values, g_{33} , at room temperature which were calculated by dividing the d_{33} values of the composites by their dielectric constant, are 48 mV m N⁻¹ and 114 mV m N⁻¹ for random and structured 20% PT-ODPA/ODA composites, respectively. The g_{33} values for random and structured 30% PT-ODPA/ODA composites are 45 mV m N⁻¹ and 90 mV m N⁻¹, respectively. The room temperature g_{33} values for the PT-ODPA/ODA composites are compared to those of PT-epoxy reference composites, as presented in our previous paper,¹³ in Fig. 7a. The DEP structured 20% PT-ODPA/ODA composites show 30% improvement in g_{33} compared to the structured 20% PT-epoxy composites, which is mainly due to their lower dielectric constant. The 20% PT-ODPA/ODA composites, with the highest g_{33} , show a tensile strength of 50 MPa and a strain at a break of 2.6%. An elastic modulus of 4.7 GPa is measured, which is in good agreement with the DMTA results. The 20% PT-ODPA/ODA flexible composite films can be folded into a bending radius as small as 2 mm without any damage.

The g_{33} values of 10%, 20% and 30% PT-ODPA/ODA composites are calculated as a function of temperature and are shown in Fig. 7b. Structured 20% PT-ODPA/ODA composites maintain their high g_{33} values of over 100 mV m N⁻¹ over the temperature range from room temperature to 175 °C. The stable g_{33} behavior of all composites over the mentioned temperature range is attributed to the thermally stable piezoelectric properties of the high T_c lead titanate,⁴⁸ as well as stable dielectric properties of ODPA/ODA, which remain nearly independent of frequency in this temperature range, as shown in Fig. 6b. Piezopolymers such as poly(vinylidene fluoride) and poly(vinylidene fluoride-co-trifluoroethylene) (P(VDF-TrFE)) show a rapid decrease above 100 °C operating temperatures.⁵⁰

4 Conclusions

This paper demonstrated the synthesis and characterization of novel high temperature piezoelectric composites by means of dielectrophoresis structuring. Fully imidized PT-ODPA/ODA composites, prepared by solution casting, show thermally stable dielectric and piezoelectric properties in the temperature range from 25 °C to 175 °C. Thermally stable functional properties of the high T_c lead titanate,⁴⁸ as well as stable dielectric properties of the fully imidized polyetherimide polymer matrix result in an improved temperature functionality of these high performance composites. The uniqueness of the presented composite manufacturing technique lies in the process design based on the molecular structure of the polymer matrix. Firstly the ODPA/ODA polymer matrix is only partially imidized to maintain relatively high dielectric permittivity and electrical conductivity, due to the polar structure, to achieve high poling efficiency. Once the composites are fully poled, the matrix is fully imidized 300 °C. Upon full imidization the imide rings completely close, resulting in a thermally stable polymer with a

very low dielectric loss. Therefore, composites of PT-PEI are shown to maintain more than 85% of room temperature piezoelectric sensitivity up to 175 °C, showing great potential for high temperature sensing and mechanical energy harvesting.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 L. Li, S. Zhang, Z. Xu, F. Wen, X. Geng, H. J. Lee and T. R. Shrout, *J. Phys. D: Appl. Phys.*, 2013, **46**, 165306.
- 2 T. Stevenson, D. G. Martin, P. I. Cowin, A. Blumfield, A. J. Bell, T. P. Comyn and P. M. Weaver, *J. Mater. Sci.: Mater. Electron.*, 2015, **26**, 9256–9267.
- 3 X. Jiang, K. Kim, S. Zhang, J. Johnson and G. Salazar, *Sensors*, 2014, **14**, 144–169.
- 4 S. Zhang and F. Yu, *J. Am. Ceram. Soc.*, 2011, **94**, 3153–3170.
- 5 I. Tumer and A. Bajwa, 35th Joint Propulsion Conference and Exhibit, Joint Propulsion Conferences, 1999.
- 6 C. Boller, *Int. J. Syst. Sci.*, 2000, **31**, 1333–1349.
- 7 Y. Sheiretov, D. Grundy, V. Zilberstein, N. Goldfine and S. Maley, *IEEE Sens. J.*, 2009, **9**, 1527–1536.
- 8 H. J. Lee, S. Zhang, Y. Bar-Cohen and S. Sherit, *Sensors*, 2014, **14**, 14526–14552.
- 9 Q. Xue, M. Stanton and G. Elfbaum, Proceedings of the IEEE Ultrasonics Symposium, 2003, pp. 1372–1375.
- 10 H. J. Lee, S. Zhang, Y. Bar-Cohen and S. Sherit, *Sensors*, 2014, **14**, 14526–14552.
- 11 F. Yu, Q. Lu, S. Zhang, H. Wang, X. Cheng and X. Zhao, *J. Mater. Chem. C*, 2015, **3**, 329–338.
- 12 D. Damjanovic, *Rep. Prog. Phys.*, 1998, **61**, 1267–1324.
- 13 H. Khanbareh, S. van der Zwaag and W. Groen, *Smart Mater. Struct.*, 2014, **23**, 105030.
- 14 H. Khanbareh, S. van der Zwaag and W. A. Groen, *J. Intell. Mater. Syst. Struct.*, 2017, DOI: 10.1177/1045389X17689928.
- 15 C. Park, Z. Ounaies, K. E. Wise and J. S. Harrison, *Polymer*, 2004, **45**, 5417–5425.
- 16 G. M. Atkinson, R. E. Pearson, Z. Ounaies, C. Park, J. S. Harrison, S. Dogan and J. A. Midkiff, TRANSDUCERS 2003 – 12th International Conference on Solid-State Sensors, Actuators and Microsystems, Digest of Technical Papers, 2003, pp. 782–785.
- 17 Z. Ounaies, C. Park, J. S. Harrison, J. G. Smith and J. Hinkley, *Proc. SPIE-Int. Soc. Opt. Eng.*, 1999, **3669**, 171–178.
- 18 Z. Ounaies, C. Park, J. S. Harrison, J. G. Smith and J. Hinkley, *Proc. SPIE-Int. Soc. Opt. Eng.*, 1999, **3669**, 171–178.

- 19 J. Simpson, Z. Ounaies and C. Fay, Materials Research Society Symposium-Proceedings, 1997, pp. 59–64.
- 20 Z. Ounaies, C. Park, J. Harrison and P. Lillehei, *J. Thermoplast. Compos. Mater.*, 2008, **21**, 393–409.
- 21 C. Park, P. O. Su, J. Su, J. G. Smith Jr. and J. S. Harrison, *Polarization Stability Of Amorphous Piezoelectric Polyimides*, 1999.
- 22 C. P. Bowen, R. E. Newnham and C. A. Randall, *J. Mater. Res.*, 1998, **13**, 205–210.
- 23 D. van den Ende, B. Bory, W. Groen and S. van der Zwaag, *J. Appl. Phys.*, 2010, **107**, 024107.
- 24 B. Jaffe, R. S. Roth and S. Marzullo, *J. Appl. Phys.*, 1954, **25**, 809–810.
- 25 R. N. R. C. Turner, P. A. Fuierer and T. Shrout, *Appl. Acoust.*, 1994, **41**, 299–324.
- 26 X. Liu, M. Zhan, Y. Shen and K. Wang, *J. Appl. Polym. Sci.*, 2011, **119**, 3253–3263.
- 27 M. Hegde, U. Lafont, B. Norder, S. J. Picken, E. T. Samulski, M. Rubinstein and T. Dingemans, *Macromolecules*, 2013, **46**, 1492–1503.
- 28 X. Zhang, B. Zhang, Y. Wu, D. Wang and T. Wang, *J. Appl. Polym. Sci.*, 2017, **134**, DOI: 10.1002/app.44889.
- 29 M. Gutiérrez, H. Khanbareh and S. van der Zwaag, *Comput. Mater. Sci.*, 2016, **112**(part A), 139–146.
- 30 B. S. Kim, S. H. Bae, Y.-H. Park and J.-H. Kim, *Macromol. Res.*, 2007, **15**, 357–362.
- 31 N. G. Devaraju, E. S. Kim and B. I. Lee, *Microelectron. Eng.*, 2005, **82**, 71–83.
- 32 S.-H. Xie, B.-K. Zhu, X.-Z. Wei, Z.-K. Xu and Y.-Y. Xu, *Composites, Part A*, 2005, **36**, 1152–1157.
- 33 L. Ding, Y. Zhang, L. Liu, J. Hu and F. Lv, *RSC Adv.*, 2016, **6**, 56812–56818.
- 34 Q. Chi, J. Sun, C. Zhang, G. Liu, J. Lin, Y. Wang, X. Wang and Q. Lei, *J. Mater. Chem. C*, 2014, **2**, 172–177.
- 35 W. Jang, D. Shin, S. Choi, S. Park and H. Han, *Polymer*, 2007, **48**, 2130–2143.
- 36 D. Wang, B. Kurish, I. Treufeld, L. Zhu and L.-S. Tan, *J. Polym. Sci., Part A: Polym. Chem.*, 2015, **53**, 422–436.
- 37 C. P. Wong, *Polymers for Electronic and Photonic Application*, Elsevier, 2013.
- 38 I. Treufeld, D. Wang, B. Kurish, L.-S. Tan and L. Zhu, *J. Mater. Chem. A*, 2014, **2**, 20683–20696.
- 39 *Broadband Dielectric Spectroscopy*, ed. F. Kremer and A. Schonhals, Springer, 2004.
- 40 D. H. Wang, J. K. Riley, S. P. Fillery, M. F. Durstock, R. A. Vaia and L.-S. Tan, *J. Polym. Sci., Part A: Polym. Chem.*, 2013, **51**, 4998–5011.
- 41 H. Khanbareh, S. van der Zwaag and W. Groen, *Smart Mater. Struct.*, 2015, **24**, 5020.
- 42 Y. T. Or, C. K. Wong, B. Ploss and F. G. Shin, *J. Appl. Phys.*, 2003, **94**, 3319–3325.
- 43 K. W. Kwok, S. T. Lau, C. K. Wong and F. G. Shin, *J. Phys. D: Appl. Phys.*, 2007, **40**, 6818–6823.
- 44 T. Yamada, T. Ueda and T. Kitayama, *J. Appl. Phys.*, 1982, **53**, 4328.
- 45 Y. Shen, Y. Lin and C.-W. Nan, *Adv. Funct. Mater.*, 2007, **17**, 2405–2410.
- 46 M. C. Araújo, C. Costa and S. Lanceros-Méndez, *J. Non-Cryst. Solids*, 2014, **387**, 6–15.
- 47 J. Yao, C. Xiong, L. Dong, C. Chen, Y. Lei, L. Chen, R. Li, Q. Zhu and X. Liu, *J. Mater. Chem.*, 2009, **19**, 2817–2821.
- 48 B. Jaffe, W. Cook Jr. and H. Jaffe, *Piezoelectric Ceramics*, Academic Press Inc., London & New York, 3rd edn, 1971.
- 49 X. Fang, X. Liu, Z.-K. Cui, J. Qian, J. Pan, X. Li and Q. Zhuang, *J. Mater. Chem. A*, 2015, **3**, 10005–10012.
- 50 C. Baur, Y. Zhou, J. Sipes, S. Priya and W. Voit, *Energy Harvesting and Systems Materials, Mechanisms, Circuits and Storage*, 2014, **1**, 167–177.